CIRCULATING LEVELS OF INTERLEUKIN-18 CORRELATE WITH SEVERITY FOLLOWING HUMAN ACUTE LIVER INJURY

Methods Analysis of SF levels in acute liver injury patients admitted to a tertiary liver center, with western blotting and immunohistochemistry for ferritin isoforms.

Results Retrospective database analysis revealed elevated admission SF (>300 μg/l) in 109/124 (87.9%) of acute liver injury patients. Extreme SF elevations (>10000 μg/l) were more common in POD (36/71, 50.7%) compared with non-POD patients (5/53, 9.4%, p<0.001). Extremely elevated admission SF was confirmed in a prospective cohort of 47 POD cases (22/47, 46.8%). In both POD cohorts, admission SF was significantly higher in patients who died/ were transplanted compared with spontaneous survivors (p=0.001, AUC 0.772 (95% CI 0.614 to 0.831) and in patients who developed hepatic encephalopathy (p=0.038) or the systemic inflammatory response syndrome (SIRS, p=0.0006). Hyperferritinaemia correlated with proinflammatory (IL-6, Spearman’s r=0.442, p=0.006; IL-8, r=0.502, p=0.001) and antiinflammatory (IL-10; r=0.349, p=0.050) cytokine release following POD, and with organ dysfunction (SOFA; r=0.529, p<0.001), but not with serum ALT (r=0.113, p=0.277). The ferritin/ALT ratio did not improve prognostic accuracy in PODs (AUC 0.706 (95% CI 0.595 to 0.817). Immunohistochemistry confirmed H and light (L) ferritin isofrm expression in both normal liver tissue and explanted tissue from ALF patients. Immunoblotting of serum from POD patients with elevated SF revealed significant amounts of circulating H-ferritin, with no circulating H-ferritin observed in healthy controls.

Conclusion Extreme elevations of SF are common following POD, and are associated with adverse outcomes. SF is a widely available biomarker that may have prognostic value in patients with POD-ALF and merits further evaluation in larger, prospective studies. The correlation with SIRS, organ failure and cytokinaemia and the observation of circulating H ferritin also suggests that SF may be a mediator of adverse outcome.

Competing interests None declared.

CIRCULATING LEVELS OF NEOPTERIN ARE ASSOCIATED WITH ADVERSE OUTCOMES FOLLOWING PARACETAMOL-INDUCED ACUTE LIVER INJURY

Methods Macrophage activation is implicated in the pathogenesis of multiorgan failure following paracetamol overdose (POD). Simple biomarkers of macrophage activation could aid earlier identification of high-risk POD patients. Neopterin is synthesised from macrophages and monocytes upon stimulation by interferon-γ and serum levels reflect the intensity of monocyte/macrophage activation.

Methods Consecutive patients (n=33, 15 (45.5%) male) admitted to the Royal Infirmary of Edinburgh with paracetamol-induced acute liver injury (ALT>1000 IU/l and coagulopathy) were enrolled. Serum neopterin levels were measured by ELISA (IBL International, Hamburg, Germany).

Results A total of 24/33 (72.7%) PODs developed hepatic encephalopathy (HE), and therefore acute liver failure. Neopterin levels were significantly higher in PODs of median 66.0 (ICR 25.4–96.6) nmol/l) compared with both chronic liver disease (10.8 (6.7–12.1) nmol/l, n=7, p<0.001) and healthy (11.4 (9.4–15.7) nmol/l, n=10, p<0.001) controls, but were similar to non-POD acute liver injury patients (52.5 (42.0–113.8) nmol/l, n=9, p>0.05). Admission neopterin levels were significantly higher in PODs who developed HE (HE (HE, 72.9 (59.5–116.7) nmol/l, n=24; no HE, 20.7 (17.5–22.1) nmol/l, n=9, p<0.0001) or the systemic inflammatory response syndrome (SIRS, 79.1 (66.7–116.7) nmol/l, p<0.0001).

Conclusion Serum neopterin is significantly higher in POD and associated with adverse outcomes. It may be a simple and sensitive biomarker for early identification and intervention in patients at highest risk of adverse outcomes following POD.

Competing interests None declared.

EXTREME HYPERFERRITINAEMIA FOLLOWING PARACETAMOL-INDUCED HUMAN ACUTE LIVER INJURY

Introduction Activated macrophages may play a critical role in the pathogenesis of acute liver failure (ALF). Serum ferritin (SF) is a circulating marker of macrophage activation, and heavy (H) isoforms of ferritin may have immunostimulatory effects. However, ferritin may be released from necrotic liver, confounding SF interpretation in ALF. The SF/ALT ratio may have prognostic value in non-paracetamol ALF, but has not been examined in patients with paracetamol (POD)-ALF.

Methods Analysis of SF levels in acute liver injury patients admitted to a tertiary liver center, with western blotting and immunohistochemistry for ferritin isoforms.
PTU-006  
THE SEQUENTIAL ORGAN FAILURE ASSESSMENT (SOFA) SCORE IS AN EFFECTIVE TRIAGE MARKER FOLLOWING STAGGERED PARACETAMOL (ACETAMINOPHEN) OVERDOSE

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Introduction The Sequential Organ Failure Assessment (SOFA) score is an effective triage marker following single time point paracetamol (acetaminophen) overdose,¹ but has not been evaluated following staggered paracetamol overdose. The aims of this study were to evaluate the prognostic accuracy of the SOFA score in a cohort of severe acute liver injury patients following staggered paracetamol overdose.

Methods Time-course analysis of 50 staggered paracetamol overdose patients admitted to a tertiary liver centre. Individual laboratory samples were correlated with the corresponding clinical parameters in relation to time from admission, and the daily SOFA score calculated.

Results A total of 39/50 (78%) patients developed hepatic encephalopathy, and therefore acute liver failure. The area under the SOFA receiver operator characteristic for death/liver transplantation was 0.87 (95% CI 75.2 to 95.7), 0.94 (95% CI 82.5 to 99.1), and 0.98 (95% CI 84.3 to 100.0) at 0, 24, and 48 h respectively post-admission. A SOFA score of <6 at tertiary care admission predicted survival with a sensitivity of 100.0% (95% CI 76.8 to 100.0) and specificity of 58.3% (95% CI 40.8 to 74.5%), compared with 85.7% (95% CI 60.6% to 97.4%) and 75.0% (95% CI 65.2% to 79.5%) respectively for the modified Kings College criteria. Only 2/21 patients with an admission SOFA score <6 required renal replacement therapy or intracerebral pressure monitoring. SOFA significantly outperformed the Model for End-stage Liver Disease at 0 (p=0.0013), 24 (p=0.0001) and 48 h (p=0.0193) following admission.

Conclusion A SOFA score <6 at tertiary care admission following a staggered paracetamol overdose carries a high negative predictive value. The SOFA score could improve triage of high risk staggered paracetamol overdose patients.

Competing interests None declared.

REFERENCE


PTU-007  
CEREBRAL OEDEMA IS RARE IN ACUTE-ON-CHRONIC LIVER FAILURE

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Introduction Acute-on-chronic liver failure (AoCLF) has a rapidly progressive disease course associated with significant mortality. Hepatic encephalopathy (HE) is common associated with hyperammonemia, systemic inflammation and hypotension. The prevalence of cerebral oedema in AoCLF is unknown. We aimed to describe the prevalence of cerebral oedema in a cohort of AoCLF adult (>18 years) patients admitted to the liver intensive treatment unit (LITU) between January 2005 and 2011.

Methods AoCLF was defined using criteria of Sarin et al.¹ Arterial ammonia (NH₃), MELD, UKELD, and organ failure (SOFA) scores were collated (results expressed as medians with ranges). Patients who had undergone cranial CT imaging were identified. Neuro-images were reported by consultant neuro-radiologists.

Results During the study period, 1008 patients with chronic liver disease (CLD) were admitted to the LITU. 173 patients (110 male) underwent neuro-imaging. Of these 81 (48 male) fulfilled criteria for AoCLF. Over the same time period 655 patients were admitted with acute liver failure. Variceal bleeding (30%) and sepsis (31%) were the most frequent precipitants of AoCLF. Compared to the CLD group, AoCLF patients were younger (50, 24–71 vs 59, 30–74, p=0.001), serum NH₃ (143, 40–305 vs 111, 28–315), grade of HE (3, 1–4 vs 1, 0–4), MELD (25, 8–40 vs 15, 6–34), SOFA (11, 2–17 vs 4, 0–14), UKELD (63, 50–75 vs 55, 44–75) and SIRS score (2, 1–3 vs 1, 0–3) were higher (p<0.001 for all). Serum sodium was lower in the AoCLF group (132 118 vs 154 136, p<0.0001). HE (≥grade 3) occurred in 66% of AoCLF patients vs 13% CLD (p<0.0001). In those with neuro-imaging, 26% were normal, 26% demonstrated increased cerebral atrophy for age, 15% small vessel disease and 10% intra-cranial haemorrhage. Cerebral oedema was seen in two patients with AoCLF, 1 post TIPSS (NH₃ 289 μmol/l) and 1 with septic shock (NH₃ 268 μmol/l). 72 patients with CLD underwent neuro-imaging with 32% showing radiological evidence of cerebral oedema. Compared to the CLD group, 30 and 90 day survival was poorer in AoCLF (52% and 42% vs 80% and 75%, log rank p<0.0001). The mode of death was that of progressive multi-organ failure (MOF). The two patients with cerebral oedema on CT suffered cerebral deaths with tonsillar herniation. AUROC analysis for survival of AoCLF identified SOFA (0.67, 95% CI 0.54 to 0.8, p=0.02) MELD (0.74, 95% CI 0.61 to 0.87, p<0.0001) and UKELD (0.87, 95% CI 0.8 to 0.97, p<0.0001).

Conclusion Our data demonstrates poor outcome in patients with AoCLF compared to those with CLD requiring admission to LITU. Mortality was attributable to MOF and although deep levels of encephalopathy requiring ventilation were common (66%), the prevalence of cerebral oedema was rare at 2%.

Competing interests None declared.

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